INSTRUCTION
for medical use of medicinal product
LEVOKLEK®

Composition:
active ingredient: chlorophyllipt extract dense;
1 suppository contains 0.05 g (50 mg) (in recalculation on dry substance) chlorophyllipt extract dense (Eucalypti) (1:15.3-10.76:1) (extractant ethanol 93-96 %);
excipients: polysorbate, hard fat

Pharmaceutical form. Suppositories.
Basic physical and chemical properties: dark green, bullet-shaped suppositories. Presence of white bloom on the surface of a suppository is allowed.

Pharmacotherapeutic group. Gynecological antiinfectives and antiseptics.
ATC Code G01A X.

Pharmacological properties.
The active ingredient of the drug is chlorophyllipt extract dense. Chlorophyllipt extract has a stimulating effect on the receptors of the mucous membranes and shows local anti-inflammatory and antiseptic activity.
The drug has antibacterial (bacteriostatic and bactericidal) activity against staphylococci.

Clinical particulars.
Indications.
Treatment of infectious and inflammatory processes of female genital organs (colpitis, vaginitis, vulvovaginitis) and intestines caused by antibiotic-resistant strains of staphylococci.

Contraindications.
Hypersensitivity to chlorophyllipt or other components of the drug.

Interaction with other medicinal products and other forms of interaction.
Data on drug interactions are not available.

Special warnings and precautions for use.
Consult a doctor before treatment.
After insertion of the first suppository wait 6-8 hours before beginning treatment.
A course of treatment is possible after confirming the absence of allergic reactions to chlorophyllipt extract spissum (probable swelling of the lips, nasal mucous membranes or throat and other allergic reactions). If any of the following symptoms occur, discontinue the drug and consult a doctor.

Use during pregnancy or lactation.
The efficacy and safety of the drug during pregnancy and lactation have not been studied, therefore the drug should not be administered to this group of patients.

Effects on ability to drive and use machines.
No data.

Posology and method of administration.
Before suppository administration it is necessary to:

– tear one suppository in the primary packaging along the perforation line of the blister pack;
– pull the edges of the film tearing it apart and releasing the suppository from the primary packaging.
The drug is intended for vaginal or rectal use.
Dose for adults: 1 suppository twice daily. The course of treatment is 10-14 days.

Paediatric patients.
There are no data on the safety and efficacy of the drug in children.

Overdose.
Major symptoms: swelling of the lips, nasal mucous membranes and throat.
Treatment: drug discontinuation. Treatment is symptomatic.

Undesirable effects.
Immune system disorders: allergic reactions, including itching, rash, flushing, angioedema (probable swelling of the lips, nasal mucous membranes, throat, etc.).

Shelf life. 2 years.

Storage. Store in original package at temperature not exceeding 25 °C. Keep away from children.

Package. 5 suppositories in a blister, 1 or 2 blisters in a pack.

Prescription status. Without prescription.

Manufacturer. Joint Stock Company «Lekhim-Kharkiv».

Manufacturer's location and place of business.
Ukraine, 61115, Kharkiv region, Kharkiv, Severyna Pototskoho street, building 36.

Date of last revision.